=> file medline biosis caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'MEDLINE' ENTERED AT 12:51:00 ON 05 SEP 2002

FILE 'BIOSIS' ENTERED AT 12:51:00 ON 05 SEP 2002 COPYRIGHT (C) 2002 BIOLOGICAL ABSTRACTS INC.(R)

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=> s reverse(w)transcript?
L1 146494 REVERSE(W) TRANSCRIPT?

=> s thermostab?
L2 31136 THERMOSTAB?

=> s 11 (9a) 12 L3 138 L1 (9A) L2

=> s 13 and (mutat? or modif? or chang? or alter?)
L4 38 L3 AND (MUTAT? OR MODIF? OR CHANG? OR ALTER?)

=> dup rem 14
PROCESSING COMPLETED FOR L4
L5 24 DUP REM L4 (14 DUPLICATES REMOVED)

=> d 1-24 ti

- L5 ANSWER 1 OF 24 CAPLUS COPYRIGHT 2002 ACS
- TI Amplifying and sequencing DNA using thermostable DNA polymerases that differentially discriminate against dideoxynucleotides and that can be differentially activated as a result of chemical modification
- L5 ANSWER 2 OF 24 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
- TI One step RT-PCR methods, enzyme mixes and kits for use in practicing the
- L5 ANSWER 3 OF 24 CAPLUS COPYRIGHT 2002 ACS
- TI Modified or mutated reverse transcriptases with high thermostability and uses thereof
- L5 ANSWER 4 OF 24 CAPLUS COPYRIGHT 2002 ACS
- TI Immunological detection of RNA: DNA hybrids on microarrays
- L5 ANSWER 5 OF 24 CAPLUS COPYRIGHT 2002 ACS
- TI Method of reversible inactivation of thermostable enzymes using chemical modification under aqueous conditions
- L5 ANSWER 6 OF 24 CAPLUS COPYRIGHT 2002 ACS
- TI Activation of 2 types of **modified** thermostable DNA polymerases at different stages in the thermo-cycler reaction for nucleic acid amplification and sequencing
- L5 ANSWER 7 OF 24 CAPLUS COPYRIGHT 2002 ACS

High temperature reverse transcription using mutant DNA polymerases TΙ DUPLICATE 2 MEDLINE ANSWER 8 OF 24 L5Differential expression of gh1 and gh2 genes by competitive rt-pcr in TΙ rainbow trout pituitary. DUPLICATE 3 ANSWER 9 OF 24 MEDLINE L5Development of a strand-specific RT-PCR based assay to detect the TТ replicative form of hepatitis C virus RNA. ANSWER 10 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 DNA polymerases from hyperthermophiles ΤI ANSWER 11 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 Thermostable DNA polymerases from Thermotoga and mutants and their use in TIDNA sequencing and amplification ANSWER 12 OF 24 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L5Hepatitis C virus in lymphoid cells of patients coinfected with human TТ immunodeficiency virus type 1: Evidence of active replication in monocytes/macrophages and lymphocytes. ANSWER 13 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 Method for reversible modification of thermostable enzymes using ТT aldehydes and its application to nucleic acid amplification ANSWER 14 OF 24 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L5 Detection for HCV with FD-thermostable reverse ΤI transcriptase mediated RT-nested PCR. ANSWER 15 OF 24 CAPLUS COPYRIGHT 2002 ACS L5Avian sarcoma-leukosis virus reverse transcriptases with improved TΙ properties for use in reverse transcription, amplification and sequencing ANSWER 16 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus TIand mutant enzymes with exonuclease activity removed ANSWER 17 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus ΤI and mutant enzymes with exonuclease activity removed DUPLICATE 4 ANSWER 18 OF 24 MEDLINE L5 Comparison of Mycobacterium 23S rRNA sequences by high-temperature reverse ΤI transcription and PCR. DUPLICATE 5 MEDLINE ANSWER 19 OF 24 L5[Use of polymerase chain reaction for determining bcr/abl mRNA in human TIchronic myeloleukemia]. Primenenie polimeraznoi tsepnoi reaktsii dlia opredeleniia bcr/abl mRNK pri khronicheskom mieloleikoze cheloveka. DUPLICATE 6 MEDLINE L5ANSWER 20 OF 24 An improved reverse transcription-polymerase chain reaction method to TΙ study apolipoprotein gene expression in Caco-2 cells. DUPLICATE 7 MEDLINE ANSWER 21 OF 24 L5 Confirmation of mutant alpha 1 Na, K-ATPase gene and transcript in Dahl TΙ salt-sensitive/JR rats. ANSWER 22 OF 24 CAPLUS COPYRIGHT 2002 ACS L5 PCR-mediated synthesis of a gene coding for the interleukin 1 receptor TΙ

antagonist

L5

ANSWER 23 OF 24

MEDLINE

```
Rapid amplification of complementary DNA from small amounts of
ΤI
    unfractionated RNA.
    ANSWER 24 OF 24 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
L5
    MODIFIED MICROMETHOD FOR DETECTING THE REVERSE TRANSCRIPTASE
ΤI
    ACTIVITY OF RETROVIRUSES IN A CULTURE MEDIUM AND IN BIOLOGICAL MATERIALS.
=> d 3 bib ab
    ANSWER 3 OF 24 CAPLUS COPYRIGHT 2002 ACS
T.5
    2001:886488 CAPLUS
AN
    136:32693
DN
    Modified or mutated reverse
TΙ
     transcriptases with high thermostability and uses
     thereof
     Smith, Michael D.; Potter, Robert Jason; Dhariwal, Gulshan; Gerard, Gary
IN
     F.; Rosenthal, Kim
     Invitrogen Corp., USA
PA
SO
     PCT Int. Appl., 103 pp.
     CODEN: PIXXD2
ידים
     Patent
    English
LΑ
FAN.CNT 1
     PATENT NO. KIND DATE
                                         APPLICATION NO. DATE
                                          _____
                     ____
     _____
     WO 2001092500 A1 20011206 WO 2001-US16861 20010525
PΙ
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                        US 2001-845157 20010501
     US 2002090618
                     A1 20020711
                          20000526
PRAI US 2000-207196P P
                    Α
     US 2001-845157
                           20010501
     US 2001-808124
                     Α
                          20010515
     The present invention provides modified reverse
AΒ
     transcriptases with increasing thermostability. The
     invention is generally related to reverse transcriptase enzymes and
     methods for the reverse transcription of nucleic acid mols., esp. mRNA
     mols. Specifically, the invention relates to reverse
     transcriptase enzymes which have been mutated or
     modified to increase thermostability, decrease terminal
     deoxynucleotidyl transferase activity, and/or increase fidelity, and to
     methods of producing, amplifying or sequencing nucleic acid mols.
     (particularly cDNA mols.) using these reverse transcriptase enzymes or
     compns. The invention also relates to nucleic acid mols. produced by
     these methods and to the use of such nucleic acid mols. to produce desired
     polypeptides. The invention also concerns kits comprising such enzymes or
     compns.
              THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 1
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
```

DUPLICATE 8

```
ANSWER 7 OF 24 CAPLUS COPYRIGHT 2002 ACS
L5
    2001:814072 CAPLUS
AN
    135:353708
DN
    High temperature reverse transcription using mutant DNA polymerases
ΤI
    Smith, Edward Soh; Elfstrom, Carita Maria; Gelfand, David Harrow; Higuchi,
IN
    Russell Gene; Myers, Thomas William; Schoenbrunner, Nancy Jeneane; Wang,
    Alice Ming
    F. Hoffmann-La Roche AG, Switz.
PA
    Eur. Pat. Appl., 23 pp.
SO
    CODEN: EPXXDW
     Patent
DT
    English
LΑ
FAN.CNT 1
                                       APPLICATION NO. DATE
                KIND DATE
     PATENT NO.
                                        _____
     EP 1152062 A2 20011107 EP 2001-109341 20010412
PΙ
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
                                         US 2001-823649
                                                         20010330
                   A1
                          20020131
     US 2002012970
                                                         20010417
                                        BR 2001-1493
                     Α
                          20011113
     BR 2001001493
                                         CN 2001-117024
                                                         20010418
     CN 1344802
                          20020417
                     Α
PRAI US 2000-198336P P
                          20000418
AB The present invention relates to improved reverse
  transcription methods using a modified
     thermostable DNA polymerases, particularly in a magnesium ion
     buffer. These methods are particularly useful in combined
     reverse-transcription/amplification reactions.
=> d 16 bib ab
     ANSWER 16 OF 24 CAPLUS COPYRIGHT 2002 ACS
L5
     1998:263203 CAPLUS
AN
     128:318803
DN
     Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus
TI
     and mutant enzymes with exonuclease activity removed
     Mamone, Joseph A.; Davis, Maria; Sha, Dan
IN
     Amersham Life Science, Inc., USA
PA
     U.S., 41 pp.
SO
     CODEN: USXXAM
DT
     Patent
     English
LΑ
FAN.CNT 1
                                        APPLICATION NO. DATE
                    KIND DATE
     PATENT NO.
                                         ______
     ______
                           19980428
                                        US 1996-766014 19961213
                     Α
     US 5744312
PΙ
     An enzymically active DNA polymerase or fragment is provided having
AΒ
     .gtoreq.80% homol. in its amino acid sequence to at least a contiguous
     40-amino-acid sequence of DNA polymerase of Thermoanaerobacter
     thermohydrosulfuricus as well as mutant enzymes where the exonuclease
     activity has been removed. Thus, deletions of up to 1/3 of the amino acid
     sequence from the N-terminus remove the exonuclease activity of the enzyme
     and are combined with a F706Y mutation to produce a thermostable
     DNA polymerase. DNA constructs derived from the full-length gene from T.
     thermohydrosulfuricus were prepd. as expression vectors for the C-terminal
     607 or 577 amino acids of the enzyme (plus an initiating Met) and a
     mutagenic oligonucleotide was designed to prep. the F412Y form of the
     588-residue mutant. The enzyme is useful for procedures requiring
     strand-displacing DNA synthesis such as strand-displacement amplification,
     for DNA sequencing, and/or for reverse transcription.
```

=> d his (FILE 'HOME' ENTERED AT 12:50:52 ON 05 SEP 2002) FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 12:51:00 ON 05 SEP 2002 146494 S REVERSE(W)TRANSCRIPT? L131136 S THERMOSTAB? 1.2 138 S L1 (9A) L2 L3 38 S L3 AND (MUTAT? OR MODIF? OR CHANG? OR ALTER?) L424 DUP REM L4 (14 DUPLICATES REMOVED)  $L_5$ => s l1 (5a) (mutat? or modif? or chang? or alter?) <----> SEARCH ENDED BY USER SEARCH ENDED BY USER => s l1 (5a) (mutat? or modif? or chang? or alter? or increas? or enhanc?) <----> User Break----> u SEARCH ENDED BY USER => s l1 (5a) (mutat? or modif? or chang? or alter? or increas? or enhanc? or improv?) (11a) thermostab? 2 FILES SEARCHED... 19 L1 (5A) (MUTAT? OR MODIF? OR CHANG? OR ALTER? OR INCREAS? OR ENHANC? OR IMPROV?) (11A) THERMOSTAB? => dup rem 16 PROCESSING COMPLETED FOR L6 16 DUP REM L6 (3 DUPLICATES REMOVED) L7 => d 1-16 ti ANSWER 1 OF 16 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE L7 Direct detection of RNA mediated by reverse transcriptase lacking RNAse H TΙ function. ANSWER 2 OF 16 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE L7 Method for improved reverse transcription at high temperatures. TΙ ANSWER 3 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Modified or mutated reverse TΙ transcriptases with high thermostability and uses thereof ANSWER 4 OF 16 CAPLUS COPYRIGHT 2002 ACS L7one step RT-PCR methods using enzyme mixes and kits comprising mutant ΤI thermostable polymerase and reverse transcriptase ANSWER 5 OF 16 CAPLUS COPYRIGHT 2002 ACS L7Activation of 2 types of modified thermostable DNA polymerases at ΤI different stages in the thermo-cycler reaction for nucleic acid amplification and sequencing ANSWER 6 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 High temperature reverse transcription using mutant DNA polymerases TIANSWER 7 OF 16 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L7

Direct detection of RNA mediated by reverse transcriptase lacking RNAse H ΤI function. ANSWER 8 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Direct detection of RNA mediated by reverse transcriptase lacking RNAse H TIANSWER 9 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Method for reversible modification of thermostable enzymes using aldehydes TΙ and its application to nucleic acid amplification ANSWER 10 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Critical factors in the preparation of representative full-length cDNA ΤI libraries. I ANSWER 11 OF 16 CAPLUS COPYRIGHT 2002 ACS ь7 Improved reverse transcription with ΤI thermostable DNA-dependent DNA polymerases in presence of betaine ANSWER 12 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Avian sarcoma-leukosis virus reverse transcriptases with improved TΤ properties for use in reverse transcription, amplification and sequencing ANSWER 13 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Chelating agents for improving thermostability of RNA in solution TIcontaining metallic ions DUPLICATE 3 MEDLINE ANSWER 14 OF 16 L7Comparison of Mycobacterium 23S rRNA sequences by high-temperature reverse ΤI transcription and PCR. ANSWER 15 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 Truncated Thermus DNA polymerases with enhanced thermostability and DNA TΙ polymerase formulations for enhancement of nucleic acid amplification ANSWER 16 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 PCR-mediated synthesis of a gene coding for the interleukin 1 receptor TΙ antagonist => d 12 bib ab ANSWER 12 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 1998:709090 CAPLUS AN DN 129:327725 Avian sarcoma-leukosis virus reverse transcriptases with improved ΤI properties for use in reverse transcription, amplification and sequencing Gerard, Gary F.; Smith, Michael D.; Chatterjee, Deb K. IN Life Technologies, Inc., USA PΑ PCT Int. Appl., 201 pp. SO CODEN: PIXXD2 Patent DTLA English FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE -----A1 19981029 WO 1998-US8072 19980422 W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG AU 1998-73601 19980422 AU 9873601 A1 19981113 20000607 EP 1998-920859 19980422 A1 EP 1005481 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO T2 20011120 JP 1998-546292 19980422 JP 2001523098 US 1999-245026 19990205 A1 20020627 US 2002081581 P 19970422 PRAI US 1997-44589P 19970617 US 1997-49874P P A3 19980422 US 1998-64057 WO 1998-US8072 W 19980422 The title reverse transcriptases comprise a mixt. of two or more proteins AΒ with reverse transcriptase activity, one or both having reduced RNase H activity, and each exhibiting a different transcription pause site. These compns. may be used for prodn. of cDNAs as well as for nucleic acid amplication and sequencing. The modified reverse transcriptases may be produced with recombinant cells. Thus, greater yields of total and full-length cDNA product using a 7.5-kb mRNA was obtained when two different RNase H- reverse transcriptases were combined than when each was used sep. in the wild-type or RNase H- form. The two reverse transcriptases used were from Rous sarcoma virus and from Moloney murine leukemia virus. It was also noted that the Rous sarcoma virus RNase Henzyme was more thermostable than the wild-type enzyme. Other expts. indicated that the combination of RNase H- .alpha. subunit with RNase H+ .beta. subunit was more thermostable than other combinations of RNase H.+-. subunits. => d 15 bib ab ANSWER 15 OF 16 CAPLUS COPYRIGHT 2002 ACS L7 1995:377249 CAPLUS AN 122:153369 DN Truncated Thermus DNA polymerases with enhanced thermostability and DNA ΤI polymerase formulations for enhancement of nucleic acid amplification Barnes, Wayne M. ΙN USA PAPCT Int. Appl., 78 pp. SO CODEN: PIXXD2 DTPatent LΑ English FAN.CNT 2 APPLICATION NO. DATE KIND DATE PATENT NO. \_\_\_\_ \_\_\_\_\_ \_\_\_\_\_ \_\_\_ WO 1994-US1867 19940222 WO 9426766 A1 19941124 PΙ W: AU, CA, JP, NZ RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE US 1993-21623 19930219 US 5436149 A 19950725 CA 1994-2156176 19940222 CA 2156176 AΑ 19941124 19940222 AU 1994-62464 A119941212 AU 9462464 B2 19960815 AU 671204 EP 1994-909742 19940222 A1 19960124 EP 693078 19990623 В1 EP 693078 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE 19940222 T2 19990216 JP 1994-522506 JP 11501801 B2 19990419 JP 2885324 AT 1994-909742 19940222 E 19990715 AT 181573 JP 1998-359199 19940222

A2 19990907

T3 19991201

ES 1994-909742

19940222

JP 11239492

PRAI US 1993-21623 A 19930219

ES 2136730

US 1994-202032 19940222 JP 1994-522506 A3 19940222 WO 1994-US1867 W 19940222

A DNA polymerase having an amino acid sequence comprising substantially AΒ the same amino acid sequence as that of Thermus aquaticus or Thermus flavus DNA polymerase, excluding the N-terminal 280 amino acid residues of Thermus aquaticus DNA polymerase or the N-terminal 279 amino acid residues of Thermus flavus DNA polymerase, and recombinant DNA sequences encoding said DNA polymerases are claimed. A formulation of thermostable or other DNA polymerases comprising a majority component comprised of at least one thermostable or other DNA polymerase of the type described above, wherein the DNA polymerase lacks 3'-exonuclease activity, and a minority component comprised of at least one thermostable DNA polymerase exhibiting 3'-exonuclease activity, and an improved method for enzymic extension of DNA strands, esp. while, but not limited to, amplifying nucleic acid sequences by polymerase chain reaction wherein the above formulation is made and used to catalyze primer extension, are also provided. Expression vector pWB254, encoding Klentaq-278 (the T. aquaticus DNA polymerase deriv.), was prepd. E. coli contg. this plasmid were used to prep. the enzyme and large-scale purifn. of the enzyme was performed. In a PCR expt., exposure to 98.degree. was not detectably detrimental to Klentaq-278. Using a 640:1 mixt. of this enzyme with Pyrococcus furiosus DNA polymerase, efficient amplification of 8.4, 12.5, 15, and 18 kb DNA fragments was demonstrated. The fidelity of the product amplified was at least equal to that obtained using P. furiosus DNA polymerase alone.

### => d 6, 11 bib ab

L7 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2002 ACS

AN 2001:814072 CAPLUS

DN 135:353708

TI High temperature reverse transcription using mutant DNA polymerases

IN Smith, Edward Soh; Elfstrom, Carita Maria; Gelfand, David Harrow; Higuchi, Russell Gene; Myers, Thomas William; Schoenbrunner, Nancy Jeneane; Wang, Alice Ming

PA F. Hoffmann-La Roche AG, Switz.

SO Eur. Pat. Appl., 23 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN. CNT 1

FAN.	PATENT NO.	KIND DATE	APPLICATION NO.	DATE
PI		A2 20011107 CH, DE, DK, ES, FR,	EP 2001-109341 GB, GR, IT, LI, LU,	20010412 , NL, SE, MC, PT,
PRAT	US 2002012970 BR 2001001493 CN 1344802	LT, LV, FI, RO A1 20020131 A 20011113 A 20020417 P 20000418	US 2001-823649 BR 2001-1493 CN 2001-117024	20010330 20010417 20010418

AB The present invention relates to improved reverse

transcription methods using a modified

thermostable DNA polymerases, particularly in a magnesium ion buffer. These methods are particularly useful in combined reverse-transcription/amplification reactions.

- L7 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2002 ACS
- AN 1998:709222 CAPLUS
- DN 129:326936

TI Improved reverse transcription with thermostable DNA-dependent DNA polymerases in presence of betaine

```
Jendrisak, Jerome J.
IN
    Epicentre Technologies Corporation, USA
PΑ
    PCT Int. Appl., 13 pp.
SO
    CODEN: PIXXD2
DT
    Patent
LΑ
    English
FAN.CNT 1
                                        APPLICATION NO. DATE
                    KIND DATE
    PATENT NO.
                                          _____
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                                                          19980415
                                        WO 1998-US7997
    WO 9848053
                     A1 19981029
PΙ
        W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
            DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
            NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
            UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,
             CM, GA, GN, ML, MR, NE, SN, TD, TG
                                        US 1997-840474
                                                           19970421
                      Α
                           20000229
     US 6030814
                                          AU 1998-71423
                                                           19980415
     AU 9871423
                      A1
                           19981113
                           20020207
     AU 743907
                      В2
                                          EP 1998-918516
                                                           19980415
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                      Α1
     EP 977891
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
                                                           19980415
                           20001017
                                          JP 1998-546266
     JP 2000513585
                      Т2
                           19970421
PRAI US 1997-840474
                      Α
                     W
                           19980415
     WO 1998-US7997
     A method of improving the synthesis of full-length cDNA transcripts by
     Mn++-dependent reverse transcriptases, preferably DNA-dependent DNA
     polymerases, is disclosed. The improvement consists in the polymn. in the
     presence of betaine.
=> d 8, 9 bib ab
     ANSWER 8 OF 16 CAPLUS COPYRIGHT 2002 ACS
L7
     1999:511279 CAPLUS
ΑN
     131:140473
DN
     Direct detection of RNA mediated by reverse transcriptase lacking RNAse H
TI
     function
     De La Rosa, Abel; Collier, Clayton D.
IN
     Digene Corporation, USA
PΑ
     PCT Int. Appl., 45 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
FAN.CNT 3
                                          APPLICATION NO. DATE
                      KIND DATE
     PATENT NO.
                                          -----
                           _____
                      ____
                                          WO 1999-US2382
                                                           19990203
     WO 9940224
                     A1
                           19990812
PΙ
         W: AU, CA
         RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
             PT, SE
                                           US 1998-20067
                                                           19980206
                            19991130
     US 5994079
                       Α
                                           CA 1999-2320102
                                                           19990203
                            19990812
                       AA
     CA 2320102
                                          AU 1999-25811
                                                           19990203
                            19990823
                       Α1
     AU 9925811
                       B2
                            20020117
     AU 742955
                                         EP 1999-905711
                                                           19990203
                            20001122
     EP 1053354
                      A1
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
                            19980206
PRAI US 1998-20067
                            19990203
     WO 1999-US2382
                       W
```

Disclosed is a method of detecting RNA mols. of interest in which reverse transcription primers unique to the RNA mol. of interest are used for reverse transcribing the RNA with a reverse transcriptase lacking RNAse H function and the resulting RNA/DNA hybrid is detected with an antibody specific for RNA/DNA hybrids. This method can be used to detect the presence of one or many specific RNA mols. which may be present in a sample, including RNA from different organisms (such as viruses, bacteria, fungi, plants, and animals), or RNA indicative of an infection, a disease state, or predisposition to a disease in an animal. The specificity of detection is increased relative to current detection methods involving probe hybridization since the reverse transcription primers are shorter and less subject to non-specific hybridization. Specificity of the disclosed method can also be increased by using a

thermostable reverse transcriptase and

performing reverse transcription at a high temp. The disclosed method can also be used to detect reverse transcriptase activity in a sample and to identify inhibitors of reverse transcriptase. Also disclosed is a method for sequencing target RNA mols. using reverse transcriptase lacking an RNAse H function. Detection of HIV-1 RNA in different samples with a 23-nucleotide biotinylated oligonucleotide as the extension primers was demonstrated.

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2002 ACS

AN 1999:783789 CAPLUS

DN 132:19613

TI Method for reversible modification of thermostable enzymes using aldehydes and its application to nucleic acid amplification

IN Ivanov, Igor; Loffert, Dirk; Kang, Jie; Ribbe, Joachim; Steinert, Kerstin

PA Qiagen G.m.b.H., Germany

SO Eur. Pat. Appl., 16 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 2

ran.		ENT	NO.		KII	ND	DATE			A	PPLI	CATI	ои ис	o.	DATE			
ΡI		9625			A2	-	1999			E	P 19	99-1	1042	б	19990	0528		
	EP	9625 R:	AT,			DE		ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
	US	6183	-	SI,	LT,		, FI, 2001			U	s 19	98-1	8395	0	1998	1031		
PRAI			-868 -183		A A		1998 1998											

OS MARPAT 132:19613

The present invention provides a method for reversible inactivation of thermostable enzymes by chem. modification under aq. conditions. This chem. modification of thermostable enzymes has surprising effects in applications in the field of mol. biol. such as nucleic acid amplification. A method for the amplification of a target nucleic acid is disclosed comprising the steps of reacting a nucleic acid with an amplification reaction mixt. and a modified thermostable enzyme, wherein said modified thermostable polymerase is prepd. by a reaction of a mixt. of a thermostable polymerase and a chem. modifying reagent. The chem. modification reagent is an aldehyde, preferably formaldehyde. Essentially complete inactivation of the enzyme at ambient temps. is achieved, with recovery of enzymic activity at temps. above 50.degree..

<sup>=&</sup>gt; s (mmlv or alv) and thermostab?
L8 9 (MMLV OR ALV) AND THERMOSTAB?

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=> dup rem 18
PROCESSING COMPLETED FOR L8
              6 DUP REM L8 (3 DUPLICATES REMOVED)
=> d 1-6 bib ab
     ANSWER 1 OF 6 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE 1
1.9
     2001:567473 BIOSIS
ΑN
     PREV200100567473
DN
     One step RT-PCR methods, enzyme mixes and kits for use in practicing the
ΤI
     Zhao, Ningyue (1); Wurst, Helmut
ΑU
     (1) Milpitas, CA USA
CS
     ASSIGNEE: Clontech Laboratories, Inc.
     US 6300073 October 09, 2001
PΙ
     Official Gazette of the United States Patent and Trademark Office Patents,
     (Oct. 9, 2001) Vol. 1251, No. 2, pp. No Pagination. e-file.
     ISSN: 0098-1133.
DT
     Patent
LA
     English
     Enzyme compositions, kits comprising the same and methods for their use in
AΒ
     one-step RT-PCR are provided. The subject enzyme compositions at least
     include a mutant thermostable DNA polymerase and a mutant
     reverse transcriptase. In preferred embodiments, the mutant
     thermostable DNA polymerase is an N-terminal deletion mutant of
     Taq polymerase and the mutant reverse transcriptase is a point mutation
     mutant of MMLV-RT. The subject kits, in addition to the above
     described mutant thermostable DNA polymerase and mutant reverse
     transcriptase, at least include one of, and usually both of, dNTPs and a
     buffer composition, where the subject kits may further include additional
     reagents, including nucleic acids, a thermostabilizing agent, a
     glycine based osmolyte and the like. In practicing the subject methods, a
     reaction mix that at least includes template RNA, the above described
     mutant polymerase and reverse transcriptase, dNTPs, buffer, and nucleic
     acid primers is prepared. The resultant reaction mixture is maintained at
     a first set of reverse transcription conditions and then a second set of
     PCR conditions, whereby amplified amounts of DNA from a template RNA(s)
     are produced.
     ANSWER 2 OF 6 CAPLUS COPYRIGHT 2002 ACS
L9
     2001:886488 CAPLUS
ΑN
     136:32693
DN
     Modified or mutated reverse transcriptases with high
ΤI
     thermostability and uses thereof
     Smith, Michael D.; Potter, Robert Jason; Dhariwal, Gulshan; Gerard, Gary
IN
     F.; Rosenthal, Kim
     Invitrogen Corp., USA
PΑ
     PCT Int. Appl., 103 pp.
SO
     CODEN: PIXXD2
     Patent
DΤ
     English
LA
FAN.CNT 1
                                           APPLICATION NO. DATE
                      KIND DATE
     PATENT NO.
     _____
                                          WO 2001-US16861 20010525
                           20011206
     WO 2001092500
                      A1
PΙ
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT,
             RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ,
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VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG A1 20020711 US 2001-845157 20010501 US 2002090618 PRAI US 2000-207196P 20000526 Ρ A 20010501 US 2001-845157 20010515 US 2001-808124 A The present invention provides modified reverse transcriptases with AB increasing thermostability. The invention is generally related of nucleic acid mols., esp. mRNA mols. Specifically, the invention

increasing thermostability. The invention is generally related to reverse transcriptase enzymes and methods for the reverse transcription of nucleic acid mols., esp. mRNA mols. Specifically, the invention relates to reverse transcriptase enzymes which have been mutated or modified to increase thermostability, decrease terminal deoxynucleotidyl transferase activity, and/or increase fidelity, and to methods of producing, amplifying or sequencing nucleic acid mols. (particularly cDNA mols.) using these reverse transcriptase enzymes or compns. The invention also relates to nucleic acid mols. produced by these methods and to the use of such nucleic acid mols. to produce desired polypeptides. The invention also concerns kits comprising such enzymes or compns.

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2002 ACS

AN 2001:573504 CAPLUS

DN 135:149586

TI Improving reverse transcription at high temperatures using thermostable CpkB Chaperonin from hyperthermophilic archaeon Pyrococcus

IN Warthoe, Peter

PA Display Systems Biotech A/s, Den.

SO U.S., 26 pp. CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

US 6271004 B1 20010807 US 2000-603185 20000626

US 1 DK 1999-897 A 19990625

PRAI DK 1999-897 A method for improved reverse transcription at high temps. is provided, wherein a thermostable chaperone protein stabilizes a reverse transcriptase, as well as reduces the RNase H activity of said reverse transcriptase. The invention further relates to a method of producing a polypeptide complex consisting of a Chaperonin and a Moloney murine leukemia virus (MMVL) reverse transcriptase, characterized by having enhanced thermostability as well as reduced RNase H activity, compared to a (MMVL) reverse transcriptase alone. The invention further relates to a kit for the prepn. of cDNA from mRNA, comprising either both stabilizing agent and reverse transcriptase or the polypeptide complex of the invention. One particular gene of interest for this invention is the gene encoding the .beta.-subunit of a mol. Chaperonin from the hyperthermophilic archaeon Pyrococcus. The present invention is related to the discovery that the CpkB polypeptide together with a reverse transcriptase generates a system having improved DNA polymerase activity at relative high temps. compared to a reverse transcriptase alone. The invention is further related to the discovery that the CpkB polypeptide inhibits the RNase H activity normally assocd. with the MMLV wild type reverse transcriptase.

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 6 CAPLUS COPYRIGHT 2002 ACS L9 1999:548573 CAPLUS AN131:282131 DN Retroviral vectors preloaded with a viral receptor-ligand bridge protein ΤI are targeted to specific cell types Boerger, Adrienne L.; Snitkovsky, Sophie; Young, John A. T. ΑU Department of Microbiology and Molecular Genetics, Harvard Medical School, CS Boston, MA, 02115, USA Proceedings of the National Academy of Sciences of the United States of SO America (1999), 96(17), 9867-9872 CODEN: PNASA6; ISSN: 0027-8424 National Academy of Sciences PΒ DTJournal English LΑ Successful targeting methods represent a major hurdle to the use of AB retroviral vectors in cell-specific gene-delivery applications. We recently described an approach for retroviral targeting with a retroviral receptor-ligand bridge protein that was bound to the cognate cell-surface ligand receptors before viral challenge. We now report a significant improvement made to this viral targeting method by using a related bridge protein, designated TVB-EGF, comprised of the extracellular domain of the TVB receptor for subgroup B avian leukosis virus fused to epidermal growth factor (EGF). The most important activity of TVB-EGF was that it allowed specific viral entry when preloaded onto virions. Furthermore, virions preloaded with TVB-EGF were thermostable and could be produced directly from virus-packaging cells. These data suggest an approach for targeting retroviral vectors to specific cell types by using virions preloaded with a retroviral receptor-ligand bridge protein and indicate that these types of bridge proteins may be useful reagents for studying the normal mechanism of retroviral entry. THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 35 ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 5 OF 6 CAPLUS COPYRIGHT 2002 ACS L9 1998:806816 CAPLUS ΑN DN 130:48291 method for highly sensitive nucleic acid detection with Imprint primers TIfor single copy detection Creighton, Steven; Gold, Larry IN Nexstar Pharmaceuticals, Inc., USA PΑ PCT Int. Appl., 54 pp. SO CODEN: PIXXD2 Patent DTEnglish LΑ FAN.CNT 1 APPLICATION NO. DATE KIND DATE PATENT NO. \_\_\_\_\_\_ \_\_\_\_\_ WO 1998-US11457 19980603 A1 19981210 WO 9855653 W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG AU 1998-78136 19980603 A1 19981221 AU 9878136 19970606 Ρ PRAI US 1997-48886P US 1998-27107 19980220 Α WO 1998-US11457 W 19980603 A novel method for the highly selective detection of a specific target

nucleic acid sequence in a sample compn. that may contain a large no. of

nucleic acids. A copy of a target nucleic acid sequence is first formed by extension from a first primer complementary to part of the target sequence. A hybrid is then formed between this copy of the target nucleic acid sequence and a second primer, and the detection of the target nucleic acid sequence is based on the formation of pyrophosphate and/or dNMP. embodiments all involve the establishment of Idling conditions using a hybrid formed between the target nucleic acid and one or more probe primer. The net result of the Idling phenomenon is the prodn. of dNMP and PPi. Imprint primers are described that synthesize a copy, or Imprint, of the target nucleic acid that highly increase the specificity of the technique. These imprint primers are wholly or partly comprised of nuclease resistant nucleic acid residues and labeled with a group such as biotin which permits subsequent attachment to a solid support. This primer is chosen so that it hybridizes to the target nucleic acid at a position that is 3' to the location of the sequences that will later be used for Idling establishment. Trapping of Imprint and elimination of non-imprint nucleic acids is performed using avidin-coated paramagnetic beads binding to biotin. The creation of a solid phase support-bound imprint can drastically reduce the complexity of the sample. Target nucleic acid detection is indicated by PPi or NADH or ATP measured in fluormetric or electrochem. or light anal. assays. The methods have the potential to detect a single copy a target nucleic acid.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 6 OF 6 MEDLINE

DUPLICATE 2

AN 89301163 MEDLINE

DN 89301163 PubMed ID: 2472758

- TI Rapid amplification of complementary DNA from small amounts of unfractionated RNA.
- AU Doherty P J; Huesca-Contreras M; Dosch H M; Pan S
- CS Department of Immunology and Rheumatology, Hospital for Sick Children, Toronto, Ontario, Canada.
- NC = GM 38420 (NIGMS)
- SO ANALYTICAL BIOCHEMISTRY, (1989 Feb 15) 177 (1) 7-10. Journal code: 0370535. ISSN: 0003-2697.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 198908
- ED Entered STN: 19900309

Last Updated on STN: 19980206 Entered Medline: 19890810

We have combined, in a rapid and straightforward manner, the synthesis and AB subsequent amplification of individual cDNA sequences from microgram quantities of unfractionated total RNA. Taql polymerase, a thermostable DNA polymerase, and Moloney murine leukemia virus ( MMLV) reverse transcriptase share similar buffer conditions; these reactions can be performed sequentially, in a single tube, without the need for purification or changes of buffer after the synthesis of cDNA. In this way, nonspecific losses of material are minimized and the required number of cells is reduced. Cell numbers, particularly from human tissues, can be limiting; the requirement for only small amounts of unfractionated RNA makes possible the isolation and characterization of cDNAs from biological materials available in limited quantities. As a demonstration system, we report the rapid synthesis and amplification of cDNA sequences corresponding to the first exon of human immunoglobulin E (IgE). MMLV reverse transcriptase is used with specific (i.e., IgE) or generic (i.e., oligo-[dT(12-18)]) oligomers to prime first strand cDNA synthesis from unfractionated RNA isolated from a human myeloma line, U-266. The necessary primers, deoxynucleotides and Taq1 polymerase,

required for second strand cDNA synthesis and the subsequent logarithmic amplification process, are then added to the reaction mixture. This technique provides a useful means of characterizing expressed and processed gene transcripts.

#### => d his (FILE 'HOME' ENTERED AT 12:50:52 ON 05 SEP 2002) FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 12:51:00 ON 05 SEP 2002 146494 S REVERSE (W) TRANSCRIPT? L131136 S THERMOSTAB? L2138 S L1 (9A) L2 L3 38 S L3 AND (MUTAT? OR MODIF? OR CHANG? OR ALTER?) L424 DUP REM L4 (14 DUPLICATES REMOVED) L5 19 S L1 (5A) (MUTAT? OR MODIF? OR CHANG? OR ALTER? OR INCREAS? OR L6 16 DUP REM L6 (3 DUPLICATES REMOVED) L7 9 S (MMLV OR ALV) AND THERMOSTAB? L8 6 DUP REM L8 (3 DUPLICATES REMOVED) 1.9 => s 12 (6a) (MMLV or ALV) L10 1 L2 (6A) (MMLV OR ALV) => d bib ab L10 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS 2001:886488 CAPLUS AN DN 136:32693 Modified or mutated reverse transcriptases with high thermostability and ΤI Smith, Michael D.; Potter, Robert Jason; Dhariwal, Gulshan; Gerard, Gary ΤN F.; Rosenthal, Kim Invitrogen Corp., USA PA PCT Int. Appl., 103 pp. SO CODEN: PIXXD2 DT Patent English LΑ FAN.CNT 1 KIND DATE APPLICATION NO. DATE PATENT NO. \_\_\_\_\_ \_\_\_\_\_ A1 20011206 WO 2001-US16861 20010525 WO 2001092500 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG A1 20020711 US 2001-845157 20010501 US 2002090618 20000526 P PRAI US 2000-207196P 20010501 US 2001-845157 Α 20010515 US 2001-808124 Α The present invention provides modified reverse transcriptases with increasing thermostability. The invention is generally related to reverse transcriptase enzymes and methods for the reverse transcription of nucleic acid mols., esp. mRNA mols. Specifically, the invention relates to

reverse transcriptase enzymes which have been mutated or modified to increase thermostability, decrease terminal deoxynucleotidyl transferase

activity, and/or increase fidelity, and to methods of producing,

amplifying or sequencing nucleic acid mols. (particularly cDNA mols.) using these reverse transcriptase enzymes or compns. The invention also relates to nucleic acid mols. produced by these methods and to the use of such nucleic acid mols. to produce desired polypeptides. The invention also concerns kits comprising such enzymes or compns.

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> dup rem 13 PROCESSING COMPLETED FOR L3 92 DUP REM L3 (46 DUPLICATES REMOVED) L11=> d 10 bib L11 ANSWER 10 OF 92 CAPLUS COPYRIGHT 2002 ACS 2001:380819 CAPLUS AN134:363664 DN Immunological detection of RNA: DNA hybrids on microarrays ΤI Lazar, James G.; Zakel, Joan M.; Strange, Christina M.; Williams, Inna R.; TN Lorincz, Attila T. Digene Corporation, USA PAPCT Int. Appl., 80 pp. SO CODEN: PIXXD2 Patent DTEnglish LA FAN.CNT 3 KIND DATE APPLICATION NO. DATE PATENT NO. \_\_\_\_\_ \_\_\_\_ \_\_\_\_\_ WO 2000-US31277 20001114 WO 2001036681 A2 20010525 PΙ A3 20011213 WO 2001036681 W: AU, BR, CA, JP RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR US 1999-440419 19991115 20010821 US 6277579 EP 2000-980379 20001114 A2 20020814 EP 1230396 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY, TR 19991115 PRAI US 1999-440419 Α US 2000-707178 20001106 Α A2 19980206 US 1998-20067 WO 2000-US31277 W 20001114

### => d 11-92 ti

- L11 ANSWER 11 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Method of reversible inactivation of thermostable enzymes using chemical modification under aqueous conditions
- L11 ANSWER 12 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Activation of 2 types of modified thermostable DNA polymerases at different stages in the thermo-cycler reaction for nucleic acid amplification and sequencing
- L11 ANSWER 13 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI High temperature reverse transcription using mutant DNA polymerases
- L11 ANSWER 14 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Method for preparing RNA reverse transcription amplification probes for microarray

DUPLICATE 4 MEDLINE L11 ANSWER 15 OF 92 Reverse transcription slippage over the mRNA secondary structure of the TΙ LIP1 gene. L11 ANSWER 16 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. Reverse transcription slippage over the mRNA secondary structure of the TILIP1 gene. DUPLICATE 5 MEDLINE L11 ANSWER 17 OF 92 Differential expression of gh1 and gh2 genes by competitive rt-pcr in rainbow trout pituitary. DUPLICATE 6 MEDLINE L11 ANSWER 18 OF 92 Development of a strand-specific RT-PCR based assay to detect the replicative form of hepatitis C virus RNA. L11 ANSWER 19 OF 92 CAPLUS COPYRIGHT 2002 ACS DNA polymerases from hyperthermophiles TIL11 ANSWER 20 OF 92 CAPLUS COPYRIGHT 2002 ACS

polymerase in the presence of magnesium

L11 ANSWER 21 OF 92 CAPLUS COPYRIGHT 2002 ACS
TI Nucleic acid ligand inhibitors of thermostable DNA polymerases, method for their selection, and their use in PCR

Reverse transcription activity from Bacillus stearothermophilus DNA

- L11 ANSWER 22 OF 92 CAPLUS COPYRIGHT 2002 ACS
  TI Thermostable DNA polymerases from Thermotoga and mutants and their use in DNA sequencing and amplification
- L11 ANSWER 23 OF 92 MEDLINE DUPLICATE 7
  TI Melanin binds reversibly to thermostable DNA polymerase and inhibits its activity.
- L11 ANSWER 24 OF 92 MEDLINE DUPLICATE 8
  TI Hepatitis C virus in lymphoid cells of patients coinfected with human immunodeficiency virus type 1: evidence of active replication in monocytes/macrophages and lymphocytes.
- L11 ANSWER 25 OF 92 MEDLINE DUPLICATE 9
  TI Quantification of porcine follicle-stimulating hormone receptor messenger ribonucleic acid by reverse transcription-competitive polymerase chain reaction.
- L11 ANSWER 26 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Direct detection of RNA mediated by reverse transcriptase lacking RNAse H function.
- L11 ANSWER 27 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Stabilization of DNA polymerases and other enzymes by cationic surfactants
- L11 ANSWER 28 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus
- L11 ANSWER 29 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Methods for DNA amplification and sequencing
- L11 ANSWER 30 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Direct detection of RNA mediated by reverse transcriptase lacking RNAse H function

L11 ANSWER 31 OF 92 CAPLUS COPYRIGHT 2002 ACS Method for reversible modification of thermostable enzymes using aldehydes TT and its application to nucleic acid amplification L11 ANSWER 32 OF 92 CAPLUS COPYRIGHT 2002 ACS Critical factors in the preparation of representative full-length cDNA ΤI libraries. I L11 ANSWER 33 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE ΤI Detection for HCV with FD-thermostable reverse transcriptase mediated RT-nested PCR. L11 ANSWER 34 OF 92 CAPLUS COPYRIGHT 2002 ACS Improved RT-PCR. One-step RT-PCR and mRNA selective PCR TΙ L11 ANSWER 35 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. Reverse transcription polymerase chain reaction method for the detection of glycopeptide resistance in enterococci. L11 ANSWER 36 OF 92 CAPLUS COPYRIGHT 2002 ACS A one-tube nucleic acid extraction and amplification (ERTPCR) method for detecting RNA viruses L11 ANSWER 37 OF 92 CAPLUS COPYRIGHT 2002 ACS High-efficiency full-length cDNA cloning TΙ L11 ANSWER 38 OF 92 CAPLUS COPYRIGHT 2002 ACS A method of cloning cell- or tissue-specific cDNAs using display of differentially expressed transcripts (DODET) L11 ANSWER 39 OF 92 CAPLUS COPYRIGHT 2002 ACS Improved reverse transcription with thermostable DNA-dependent DNA polymerases in presence of betaine L11 ANSWER 40 OF 92 CAPLUS COPYRIGHT 2002 ACS Avian sarcoma-leukosis virus reverse transcriptases with improved properties for use in reverse transcription, amplification and sequencing L11 ANSWER 41 OF 92 CAPLUS COPYRIGHT 2002 ACS Sulfates and acetates for relief of reverse transcriptase inhibition of reverse transcriptase-polymerase chain reaction L11 ANSWER 42 OF 92 CAPLUS COPYRIGHT 2002 ACS Thermostable DNA polymerase from Carboxydothermus hydrogenoformans L11 ANSWER 43 OF 92 CAPLUS COPYRIGHT 2002 ACS Thermostable DNA polymerase from Anaerocellum thermophilum ТΤ L11 ANSWER 44 OF 92 CAPLUS COPYRIGHT 2002 ACS Endogenous ribonuclease inhibitors of mammals, cDNAs encoding them, and ΤT their uses L11 ANSWER 45 OF 92 CAPLUS COPYRIGHT 2002 ACS Nucleic acid ligand inhibitors to DNA polymerases L11 ANSWER 46 OF 92 CAPLUS COPYRIGHT 2002 ACS Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus and mutant enzymes with exonuclease activity removed L11 ANSWER 47 OF 92 CAPLUS COPYRIGHT 2002 ACS Cloning and gene sequence of a thermostable DNA polymerase from Bacillus TI

pallidus and its use for strand displacement amplification L11 ANSWER 48 OF 92 CAPLUS COPYRIGHT 2002 ACS Chelating agents for improving thermostability of RNA in solution TТ containing metallic ions L11 ANSWER 49 OF 92 CAPLUS COPYRIGHT 2002 ACS RT-PCR for DNA amplification using thermostable RNase H to improve amplification efficiency and detection sensitivity L11 ANSWER 50 OF 92 CAPLUS COPYRIGHT 2002 ACS cloning, sequence, and expression of a thermostable DNA polymerase gene ΤI from Bacillus pallidus L11 ANSWER 51 OF 92 CAPLUS COPYRIGHT 2002 ACS Detection of hepatitis G virus replication sites by using highly strand-specific Tth-based reverse transcriptase PCR DUPLICATE 11 L11 ANSWER 52 OF 92 MEDLINE Recombinant His-tagged DNA polymerase. I. Cloning, purification and partial characterization of Thermus thermophilus recombinant DNA polymerase. L11 ANSWER 53 OF 92 CAPLUS COPYRIGHT 2002 ACS Thermostabilization and thermoactivation of thermolabile enzymes by TТ trehalose and its application for the synthesis of full length cDNA L11 ANSWER 54 OF 92 CAPLUS COPYRIGHT 2002 ACS Tertiary structure model of FD-thermostable reverse TΙ transcriptase (FD-TRT) and its structure-based homology analysis L11 ANSWER 55 OF 92 CAPLUS COPYRIGHT 2002 ACS Characterization of FD-thermostable reverse тт transcriptase (FD-TRT) L11 ANSWER 56 OF 92 CAPLUS COPYRIGHT 2002 ACS Partial enzymic characteristics of FD thermostable reverse transcriptase (FD-TRT) DUPLICATE 12 L11 ANSWER 57 OF 92 MEDLINE Differential display with carboxy-X-rhodamine-labeled primers and the selection of differentially amplified cDNA fragments without cloning. L11 ANSWER 58 OF 92 CAPLUS COPYRIGHT 2002 ACS Thermostable DNA polymerase from Thermoanaerobacter thermohydrosulfuricus TТ and mutant enzymes with exonuclease activity removed L11 ANSWER 59 OF 92 CAPLUS COPYRIGHT 2002 ACS Use of manganese, metal ion buffer, and thermostable DNA polymerase for coupled high temperature reverse transcription and polymerase chain reaction. L11 ANSWER 60 OF 92 CAPLUS COPYRIGHT 2002 ACS Encapsulation of thermostable enzymes in heat-labile wax beads or ΤI liposomes for release upon heating DUPLICATE 13 L11 ANSWER 61 OF 92 MEDLINE A simple reverse transcription-polymerase chain reaction for dengue type 2 virus identification. DUPLICATE 14 L11 ANSWER 62 OF 92 MEDLINE The use of the reverse transcription-competitive polymerase chain reaction to investigate the in vivo regulation of gene expression in small tissue samples.

L11 ANSWER 63 OF 92 MEDLINE DUPLICATE 15

- TI Detection and identification of dengue virus isolates from Brazil by a simplified reverse transcription-polymerase chain reaction (RT-PCR) method.
- L11 ANSWER 64 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Methods for reverse transcription using thermostable DNA polymerase to amplify and detect target RNA
- L11 ANSWER 65 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE 16
- TI RT-PCR-based genotyping for swine major histocompatibility complex (SLA) class II genes.
- L11 ANSWER 66 OF 92 MEDLINE DUPLICATE 17
- TI Phylogenetic footprinting of the human cytochrome c oxidase subunit VB promoter.
- L11 ANSWER 67 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI a thermostable nucleic acid polymerase from Thermus sps17 for use in nucleic acid amplification and the gene encoding it
- L11 ANSWER 68 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Use of manganese, metal ion buffer, and thermostable DNA polymerase for coupled high temperature reverse transcription and polymerase chain reaction.
- L11 ANSWER 69 OF 92 MEDLINE DUPLICATE 18
- TI [Use of thermostable DNA polymerase from Thermus thermophilus KTP in a combined reverse transcription and amplification reaction for detecting CD4 receptor mRNA]. Ispol'zovanie termostabil'noi DNK-polimerazy iz Thermus thermophilus KTP v sovmeshchennoi reaktsii obratnoi transkriptsii i amplifikatsii dlia detektsii mRNK retseptora CD-4.
- L11 ANSWER 70 OF 92 MEDLINE DUPLICATE 19
- II [Use of thermostable DNA polymerase from Thermus thermophilus KTP in a combined reverse transcription and amplification reaction of detecting interleukin 2alpha RNA and determining expression of the multidrug resistance gene (MDR-1)].

  Ispol'zovanie termostabil'noi DNK-polimerazy iz Thermus thermophilus STP v sovmeshchennoi reaktsii obratnoi transkriptsii i amplifikatsii dlia detektsii RNK interleikina 2alpha i opredelenie ekspressii gena mnozhestvennoi lekarstvennoi ustichivosti (MDR-1).
- L11 ANSWER 71 OF 92 MEDLINE DUPLICATE 20
- TI Comparison of Mycobacterium 23S rRNA sequences by high-temperature reverse transcription and PCR.
- L11 ANSWER 72 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Use of PCR in detection of antisense transcripts in HTLV-I-infected patients and human T-cell lines
- L11 ANSWER 73 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Truncated Thermus DNA polymerases with enhanced thermostability and DNA polymerase formulations for enhancement of nucleic acid amplification
- L11 ANSWER 74 OF 92 MEDLINE DUPLICATE 21
- TI [Use of polymerase chain reaction for determining bcr/abl mRNA in human

chronic myeloleukemia]. Primenenie polimeraznoi tsepnoi reaktsii dlia opredeleniia bcr/abl mRNK pri khronicheskom mieloleikoze cheloveka.

- L11 ANSWER 75 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Detection of mRNA expression in a single cell by direct RT-PCR
- L11 ANSWER 76 OF 92 MEDLINE DUPLICATE 22
- TI Demonstration of in vitro infection of chimpanzee hepatocytes with hepatitis C virus using strand-specific RT/PCR.
- L11 ANSWER 77 OF 92 MEDLINE DUPLICATE 23
- TI An improved reverse transcription-polymerase chain reaction method to study apolipoprotein gene expression in Caco-2 cells.
- L11 ANSWER 78 OF 92 MEDLINE DUPLICATE 24
- TI Separate detection of the two complementary RNA strands of hepatitis A virus.
- L11 ANSWER 79 OF 92 MEDLINE DUPLICATE 25
- TI Confirmation of mutant alpha 1 Na, K-ATPase gene and transcript in Dahl salt-sensitive/JR rats.
- L11 ANSWER 80 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Single-step amplification method for RNA
- L11 ANSWER 81 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Efficient extraction of viral RNA for PCR amplification
- L11 ANSWER 82 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI PCR-mediated synthesis of a gene coding for the interleukin 1 receptor antagonist
- L11 ANSWER 83 OF 92 MEDLINE DUPLICATE 26
- TI Molecular cloning of a mouse extracellular signal regulated kinase (erk-1).
- L11 ANSWER 84 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI A thermostable nucleic acid polymerase purified from Thermosipho africanus cloning of the gene
- L11 ANSWER 85 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI RNA detection by polymerase chain reaction
- L11 ANSWER 86 OF 92 MEDLINE DUPLICATE 27
- TI Improved detection of hepatitis C virus RNA by reverse transcription and polymerase chain reaction.
- L11 ANSWER 87 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Rapid and contamination-safe nested PCR as a one-tube-reaction with thermostable RTTH-reverse-transcriptase /polymerase and CG-clamp primers.
- L11 ANSWER 88 OF 92 CAPLUS COPYRIGHT 2002 ACS
- TI Reverse transcription using thermostable DNA polymerases
- L11 ANSWER 89 OF 92 MEDLINE DUPLICATE 28
- TI Reverse transcription and DNA amplification by a Thermus thermophilus DNA polymerase.
- L11 ANSWER 90 OF 92 MEDLINE

- Rapid amplification of complementary DNA from small amounts of ΤI unfractionated RNA.
- L11 ANSWER 91 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- MODIFIED MICROMETHOD FOR DETECTING THE REVERSE TRANSCRIPTASE ACTIVITY OF RETROVIRUSES IN A CULTURE MEDIUM AND IN BIOLOGICAL MATERIALS.
- L11 ANSWER 92 OF 92 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- ENZYMATIC SYNTHESIS AND CHARACTERIZATION OF DNA COMPLEMENTARY TO ΤI CERULOPLASMIN MESSENGER RNA FROM RAT LIVER.

### => d 88 bib ab

L11 ANSWER 88 OF 92 CAPLUS COPYRIGHT 2002 ACS

1991:552514 CAPLUS AN

115:152514 DN

Reverse transcription using thermostable DNA  $\mathtt{TI}$ polymerases

Gelfand, David H.; Myers, Thomas W. IN

US 1990-609157 B2 19901102

Cetus Corp., USA PΑ

PCT Int. Appl., 52 pp. SO

CODEN: PIXXD2

Patent DT

English LΑ

FAN. CNT 27

FAN.		27 TENT NO.	MIND	ኮአጥሮ		APPLICATION NO.	DATE
	PA.	ENT NO.	KIND				
ΡI	WO	9109944	A2	19910711		WO 1990-US7641	19901221
			A3				
		W: AU, CA,	JP, US				
		RW: AT, BE,	CH, DE	, DK, ES,	FR,	GB, GR, IT, LU, NL	, SE
	US	5322770		19940621		US 1989-455611	19891222
	CA	2071213				CA 1990-2071213	
	ΑU	9172444	A1	19910724		AU 1991-72444	19901221
	ΑU	656315		19950202			
	ΕP	506889	A1	19921007		EP 1991-904087	19901221
	ΕP	506889	B1				
						GB, GR, IT, LI, LU	, NL, SE
	JΡ	05505105	<b>T</b> 2	19930805		JP 1991-504344	
	ΑT	151112	E	19970415		AT 1991-904087	
	ES	2100945	Т3	19970701		ES 1991-904087	
		09224682	A2			JP 1996-246648	
		2968585	B2	19991025		JP 1990-504344	
		5407800	А	19950418		US 1993-80243	19930617
		5618703	A	19970408		US 1994-199509	19940222
		5641864	Α	19970624		US 1994-311612	19940922 19950206
		5618711	Α	19970408		US 1995-384490 US 1995-459383	19950200
		5789224		19980804		US 1995-459383	19950602
		5795762	A			US 1995-458819	19930002
PRAI		1989-455611		19891222			
		1989-455967	A	19891222			
		1990-585471	A2	19900920			
		1986-899241	B2	19860822			
		1987-63509	A2	19870617 19880112			
		1988-143441	B2	19900515			
		1990-523394 1990-557517	A2 B2	19900313			
		1990-590213	В2 В2	19900724			
		1990-590213	A2	19900928			
		1990-590490	B2	19900928			
		1990-390490	D2	10001100			

JΡ	1991-502929	A3	19901221
WO	1990-US7641	Α	19901221
US	1991-746121	В1	19910815
US	1992-880478	B1	19920506
US	1993-977434	A1	19930223
US	1993-82182	A1	19930624
US	1993-148133	В1	19931102
US	1994-199509	A1	19940222
US	1995-384490	A3	19950206

AB A method for reverse transcription of RNA using the heat-stable DNA polymerases of Thermus and without use of reverse transcriptase is described. Optimization expts. and methods for direct amplification of the cDNA are reported.

### => d 54-56 bib ab

- L11 ANSWER 54 OF 92 CAPLUS COPYRIGHT 2002 ACS
- AN 1998:632665 CAPLUS
- DN 130:22145
- TI Tertiary structure model of FD-thermostable reverse transcriptase (FD-TRT) and its structure-based homology analysis
- AU Zhang, Kun; Wang, Shunde; Zheng, Zuohua; Mao, Yumin
- CS Department Physiology Biophysics, Fudan University, Shanghai, 200433, Peop. Rep. China
- SO Fudan Xuebao, Ziran Kexueban (1998), 37(4), 455-461 CODEN: FHPTAY; ISSN: 0427-7104
- PB Shanghai Kexue Jishu Chubanshe
- DT Journal
- LA Chinese
- Using automatic homol. modeling methods and taking the crystal structure of Taq polymerase as model block, the authors adopt a combined method to build a tertiary structure model of FD-thermostable reverse transcriptase (FD-TRT). The model makes them possible to investigate the structure basis for the functional difference between FD-TRT and other proteins in the DNA polymerase family.
  - Functional sites of the reverse transcriptase are discussed.
- L11 ANSWER 55 OF 92 CAPLUS COPYRIGHT 2002 ACS
- AN 1998:437137 CAPLUS
- DN 129:199637
- TI Characterization of FD-thermostable reverse transcriptase (FD-TRT)
- AU Yin, Changchuan; Yan, Xuehen; Zheng, Zuohua; Huang, Xiaoyu; Mao, Yumin
- CS State Key Laboratory of Genetic Engineering, Fudan University, Shanghai, Peop. Rep. China
- SO Fudan Xuebao, Ziran Kexueban (1998), 37(2), 225-228 CODEN: FHPTAY; ISSN: 0427-7104
- PB Shanghai Kexue Jishu Chubanshe
- DT Journal
- LA Chinese
- AB FD thermostable reverse transcriptase (FD-TRT) was isolated from a Thermus strain. An optimal assay method of FD-TRT was developed using the yeast rRNA as template. FD-TRT showed optimal activity on the reaction condition of 25 mmol/L Tris-HCl (pH 8.5, 25.degree.C), 25 mmol/L (NH4)2SO4, 2 mmol/L MnCl2, 100 .mu.g/mL gelatin, 5 unit RNasin, 250 .mu.mol/L each of four dNTPs, 1 .mu.Ci 3H-dCTP, 12 .mu.g RNA, and 25 pmol primers. The activity ratios of reverse transcriptase to DNA polymerase were 0.056 and 0.0045 for FD-TRT and Taq DNA polymerase, resp.
- L11 ANSWER 56 OF 92 CAPLUS COPYRIGHT 2002 ACS

AN 1998:321879 CAPLUS

DN 129:64711

TI Partial enzymic characteristics of FD thermostable reverse transcriptase (FD-TRT)

- AU Zheng, Zuo-Hua; Zhou, Zong-Xiang; Yin, Chang-Chuan; Ji, Chao-Neng; Mao, Yu-Min
- CS Inst. of Genetics, Fudan Univ., Shanghai, 200433, Peop. Rep. China
- SO Zhongguo Shengwu Huaxue Yu Fenzi Shengwu Xuebao (1998), 14(2), 170-174 CODEN: ZSHXF2; ISSN: 1007-7626
- PB Zhongguo Shengwu Huaxue Yu Fenzi Shengwu Xuebao Bianweihui
- DT Journal
- LA Chinese
- AB FD thermostable reverse transcriptase

(FD-TRT) was isolated from a Thermus strain. Some of its enzymic properties were studied through RT-PCR method. FD-TRT can endure 95.degree.C, and its optimal reaction temp. is around 65-70.degree.C when most of the coiled structure of RNA are opened, thus the high temp. can improve the efficiency of reverse transcription. Also as the specificity of recognition between primer and template is increased, it will improve the specificity of reverse transcription. The optimal reaction condition of FD-TRT is as follows: 25 mmol/L Tris-HCl (pH 8.8), 15 mmol/L (NH4)2SO4, 100 .mu.g/mL gelatin, 500 .mu.mol/L dNTPs, 25 pmol reverse transcription primer, 1 mmol/L Mn-Cl2, 2 U FD-TRT, incubation at 65-70.degree.C, .alpha. globin mRNA can be efficiently detected from less than 5 pg total RNA of human peripheral blood cell with RT-PCR conducted by FD-TRT under the above condition.

### => d 22, 30 bib ab

L11 ANSWER 22 OF 92 CAPLUS COPYRIGHT 2002 ACS

AN 2000:46954 CAPLUS

DN 132:103728

TI Thermostable DNA polymerases from Thermotoga and mutants and their use in DNA sequencing and amplification

IN Hughes, A. John; Chatterjee, Deb K.

PA Life Technologies, Inc., USA

SO U.S., 65 pp., Cont.-in-part of U. S. Ser. No. 689,818, abandoned. CODEN: USXXAM

DT Patent

LA English

FAN. CNT 6

PAN.	CMI 6				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
			<del>_</del>		
ΡI	US 6015668	Α	20000118	US 1996-706706	19960906
	US 5912155	Α	19990615	us 1995-370190	19950109
	US 5939301	Α	19990817	US 1995-537400	19951002
PRAI	US 1994-316423	В2	19940930		
	US 1995-370190	A2	19950109		
	บร 1995-525057	В2	19950908		
	US 1995-537397	B1	19951002		
	US 1995-537400	A2	19951002		
	us 1995-576759	A2	19951221		
	US 1996-689818	B2	19960814		
70.00	The method of st	nthesi	zina sequencia	ng, and amplifying	a double

The method of synthesizing, sequencing, and amplifying a double strand DNA using the Thermotoga DNA polymerase and the kit required are disclosed. The invention relates to a thermostable DNA polymerase from Thermotoga neapolitana (Tne) and mutants. The mutant DNA polymerase has at least one mutation selected from the group consisting of (1) a first mutation that substantially reduces or eliminates 3'.fwdarw.5' exonuclease activity of said DNA polymerase; (2) a second mutation that substantially reduces or eliminates 5'.fwdarw.3' exonuclease activity of said DNA polymerase; (3) a

third mutation in the O helix of said DNA polymerase resulting in said DNA polymerase becoming non-discriminating against dideoxynucleotides. The present invention also relates to the cloning and expression of the wild type or mutant DNA polymerases in E. coli, to DNA mols. contg. the cloned gene, and to host cells which express said genes.

RE.CNT 70 THERE ARE 70 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 30 OF 92 CAPLUS COPYRIGHT 2002 ACS

AN 1999:511279 CAPLUS

DN 131:140473

TI Direct detection of RNA mediated by reverse transcriptase lacking RNAse H function

IN De La Rosa, Abel; Collier, Clayton D.

PA Digene Corporation, USA

SO PCT Int. Appl., 45 pp.

CODEN: PIXXD2

DT Patent

LA English

באאו האות 3

FAN.	CNT	3																
	PAT	CENT	NO.		KI	ND	DATE			Al	PLIC	CATI	ои ис	ο.	DATE			
PI	WO	9940			<b>-</b>	1	1999	0812		WC	199	99-U	s2382	2	1999	0203		
			AU, AT, PT,	BE,	CH,	CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,
	US	5994	•	מט	А		1999	1130		U.	3 19	98-2	0067		1998	0206		
	CA	2320	102		A	A	1999	0812		C.	199	99-2	3201	02	1999			
	AU	9925	811		Α	1	1999	0823		ΑŪ	J 19	99-2	5811		1999	0203		
	ΑU	7429	55		В	2	2002	0117					_					
	ΕP	1053	354		Α	1	2000	1122					0571	_	1999			
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
			ΙE,	FI														
PRAI	US	1998	-200	67	Α		1998	0206										
	WO	1999	-US2	382	W		1999	0203										

Disclosed is a method of detecting RNA mols. of interest in which reverse transcription primers unique to the RNA mol. of interest are used for reverse transcribing the RNA with a reverse transcriptase lacking RNAse H function and the resulting RNA/DNA hybrid is detected with an antibody specific for RNA/DNA hybrids. This method can be used to detect the presence of one or many specific RNA mols. which may be present in a sample, including RNA from different organisms (such as viruses, bacteria, fungi, plants, and animals), or RNA indicative of an infection, a disease state, or predisposition to a disease in an animal. The specificity of detection is increased relative to current detection methods involving probe hybridization since the reverse transcription primers are shorter and less subject to non-specific hybridization. Specificity of the disclosed method can also be increased by using a thermostable

## reverse transcriptase and performing reverse

transcription at a high temp. The disclosed method can also be used to detect reverse transcriptase activity in a sample and to identify inhibitors of reverse transcriptase. Also disclosed is a method for sequencing target RNA mols. using reverse transcriptase lacking an RNAse H function. Detection of HIV-1 RNA in different samples with a 23-nucleotide biotinylated oligonucleotide as the extension primers was demonstrated.

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

# (FILE 'HOME' ENTERED AT 12:50:52 ON 05 SEP 2002)

	FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 12:51:00 ON 05 SEP 2002
L1	146494 S REVERSE (W) TRANSCRIPT?
L2	31136 S THERMOSTAB?
L3	138 S L1 (9A) L2
L4	38 S L3 AND (MUTAT? OR MODIF? OR CHANG? OR ALTER?)
L5	24 DUP REM L4 (14 DUPLICATES REMOVED)
L6	19 S L1 (5A) (MUTAT? OR MODIF? OR CHANG? OR ALTER? OR INCREAS? OR
L7	16 DUP REM L6 (3 DUPLICATES REMOVED)
L8	9 S (MMLV OR ALV) AND THERMOSTAB?
L9	6 DUP REM L8 (3 DUPLICATES REMOVED)
L10	1 S L2 (6A) (MMLV OR ALV)

=> file medline biosis caplus COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FILE 'MEDLINE' ENTERED AT 07:11:36 ON 10 SEP 2002

FILE 'BIOSIS' ENTERED AT 07:11:36 ON 10 SEP 2002 COPYRIGHT (C) 2002 BIOLOGICAL ABSTRACTS INC.(R)

FILE 'CAPLUS' ENTERED AT 07:11:36 ON 10 SEP 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS)

=> s (divers? or variab?) (5a) (mmlv) (3a) reverse(w)transcriptase#
L1 0 (DIVERS? OR VARIAB?) (5A) (MMLV) (3A) REVERSE(W) TRANSCRIPTASE#

=> s (divers? or variab?) and (mmlv) (3a) reverse(w)transcriptase#
L2 0 (DIVERS? OR VARIAB?) AND (MMLV) (3A) REVERSE(W) TRANSCRIPTASE#

=> s (divers? or variab?) and reverse(w)transcriptase#
L3 4640 (DIVERS? OR VARIAB?) AND REVERSE(W) TRANSCRIPTASE#

=> s (divers? or variab?) (7a) reverse(w)transcriptase#
L4 112 (DIVERS? OR VARIAB?) (7A) REVERSE(W) TRANSCRIPTASE#

=> s 14 and mmlv L5 0 L4 AND MMLV

=> dup rem 14
PROCESSING COMPLETED FOR L4
L7 57 DUP REM L4 (55 DUPLICATES REMOVED)

=> d 1-57 ti

- L7 ANSWER 1 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- Genetic diversity of protease and reverse
  transcriptase sequences in non-subtype-B human immunodeficiency
  virus type 1 strains: Evidence of many minor drug resistance mutations in
  treatment-naive patients.
- L7 ANSWER 2 OF 57 MEDLINE DUPLICATE 1
- TI Isotype-switched immunoglobulin genes with a high load of somatic hypermutation and lack of ongoing mutational activity are prevalent in mediastinal B-cell lymphoma.
- L7 ANSWER 3 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Diversity, origin, and distribution of retrotransposons (gypsy and copia) in conifers.
- L7 ANSWER 4 OF 57 CAPLUS COPYRIGHT 2002 ACS
- TI Targeting human immunodeficiency virus type 1 reverse transcriptase by intracellular expression of single-chain variable fragments to inhibit early stages of the viral life cycle. [Erratum to document cited in CA124:340469]
- L7 ANSWER 5 OF 57 MEDLINE

DUPLICATE 2

TI Possible regulation of telomerase activity by transcription and alternative splicing of telomerase reverse transcriptase in human melanoma.

L7 ANSWER 6 OF 57 MEDLINE DUPLICATE 3

- TI Somatostatin induces migration of acute myeloid leukemia cells via activation of somatostatin receptor subtype 2.
- L7 ANSWER 7 OF 57 MEDLINE DUPLICATE 4
- TI Human immunodeficiency virus type 1 protease genotype predicts immune and viral responses to combination therapy with protease inhibitors (PIs) in PI-naive patients.
- L7 ANSWER 8 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Isotype-switched immunoglobulin genes with a high load of somatic hypermutation and lack of ongoing mutational activity are prevalent in mediastinal B cell lymphoma.
- L7 ANSWER 9 OF 57 CAPLUS COPYRIGHT 2002 ACS
- TI Effects of HIV-1 clade diversity on HIV-1 virulence and antiretroviral drug sensitivity
- L7 ANSWER 10 OF 57 CAPLUS COPYRIGHT 2002 ACS
- TI Pathogenicity and DNA sequence of variable region of VP2 gene of cell-adapted strain X of infectious bursal disease virus
- L7 ANSWER 11 OF 57 MEDLINE DUPLICATE 5
- Genetic diversity of protease and reverse
  transcriptase sequences in non-subtype-B human immunodeficiency
  virus type 1 strains: evidence of many minor drug resistance mutations in
  treatment-naive patients.
- L7 ANSWER 12 OF 57 MEDLINE DUPLICATE 6
- TI Analytical variables of reverse transcription-polymerase chain reaction-based detection of disseminated prostate cancer cells.
- L7 ANSWER 13 OF 57 CAPLUS COPYRIGHT 2002 ACS
- TI Partial Molecular Alignment via Local Structure Analysis
- L7 ANSWER 14 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE 7
- TI Expression of Trigonopsis variabilis D-amino acid oxidase gene in Escherichia coli and characterization of its inactive mutants.
- L7 ANSWER 15 OF 57 MEDLINE DUPLICATE 8
- TI Functional and genetic integrity of the CD8 T-cell repertoire in advanced HIV infection.
- L7 ANSWER 16 OF 57 MEDLINE DUPLICATE 9
- TI Sequence diversity of the reverse transcriptase of human immunodeficiency virus type 1 from untreated Brazilian individuals.
- L7 ANSWER 17 OF 57 MEDLINE DUPLICATE 10
- TI Telomerase and the maintenance of chromosome ends.
- L7 ANSWER 18 OF 57 MEDLINE DUPLICATE 11
- TI The impact of biochemical methods for single muscle fibre analysis.
- L7 ANSWER 19 OF 57 MEDLINE
- TI Oligoclonal expansions of T-cell repertoire in gastric mucosa associated lymphoid tissue type B-cell lymphoma and adjacent gastritis.

DUPLICATE 12 ANSWER 20 OF 57 MEDLINE L7 Antiendotoxin agents share molecular homology within their TΙ lipopolysaccharide binding domains. DUPLICATE 13 ANSWER 21 OF 57 MEDLINE L7 A molecular-field-based similarity study of non-nucleoside HIV-1 reverse ΤI transcriptase inhibitors. DUPLICATE 14 ANSWER 22 OF 57 MEDLINE L7 Haemophilus ducreyi secretes a filamentous hemagglutinin-like protein. TΙ ANSWER 23 OF 57 CAPLUS COPYRIGHT 2002 ACS L7 Variability in repeated consecutive measurements of plasma human TТ immunodeficiency virus RNA in persons receiving stable nucleoside reverse transcriptase inhibitor therapy or no treatment DUPLICATE 15 ANSWER 24 OF 57 MEDLINE L7 Consensus-degenerate hybrid oligonucleotide primers for amplification of TΙ distantly related sequences. DUPLICATE 16 ANSWER 25 OF 57 MEDLINE L7 Possible roles of nucleocapsid protein of MoMuLV in the specificity of ΤI proviral DNA synthesis and in the genetic variability of the virus. ANSWER 26 OF 57 CAPLUS COPYRIGHT 2002 ACS L7 A quantum molecular similarity approach to anti-HIV activity ITMEDLINE L7 ANSWER 27 OF 57 Gene expression of malignant rhabdoid tumor cell lines by reverse ΤI transcriptase-polymerase chain reaction. DUPLICATE 17 ANSWER 28 OF 57 MEDLINE L7Structural variation among retroviral primer-DNA junctions: solution TΤ structure of the HIV-1 (-)-strand Okazaki fragment r(gcca)d(CTGC).d(GCAGTGGC). DUPLICATE 18 ANSWER 29 OF 57 MEDLINE L7 Preparation of an antifibrin thrombus-specific murine/human chimeric TImonoclonal antibody Fab fragment in Escherichia coli. MEDLINE ANSWER 30 OF 57 ь7 Evidence of a butterfly-like configuration of structurally diverse ΤI allosteric inhibitors of the HIV-1 reverse transcriptase DUPLICATE 20 ANSWER 31 OF 57 MEDLINE L7T cell receptor clonal diversity following allogeneic marrow grafting. ΤI DUPLICATE 21 ANSWER 32 OF 57 MEDLINE L7 Assessment of a standardized reverse-transcriptase PCR assay for TIquantifying HIV-1 RNA in plasma and serum. MEDLINE ANSWER 33 OF 57 L7 Preparation of samples for polymerase chain reaction in situ. ΤI DUPLICATE 23 MEDLINE ANSWER 34 OF 57 L7 HIV as the cause of AIDS. ΤI DUPLICATE 24 ANSWER 35 OF 57 MEDLINE ь7 Multiple cysteine proteinases of the pathogenic protozoon Tritrichomonas ΤI foetus: identification of seven diverse and differentially expressed

MEDLINE Oligoclonal expansion of V delta 1+ gamma/delta T-cells in systemic sclerosis patients. MEDLINE

- DUPLICATE 25 ANSWER 37 OF 57 L7
- Comparative anti-HIV evaluation of diverse HIV-1-specific ТT reverse transcriptase inhibitor-resistant virus isolates demonstrates the existence of distinct phenotypic subgroups.
- DUPLICATE 26 ANSWER 38 OF 57 MEDLINE L7
- Phylogenetic comparison of retron elements among the myxobacteria: TТ evidence for vertical inheritance.
- DUPLICATE 27 ANSWER 39 OF 57 MEDLINE L7
- Kinetic and mutational analysis of human immunodeficiency virus type 1TIreverse transcriptase inhibition by inophyllums, a novel class of non-nucleoside inhibitors.
- **DUPLICATE 28** ANSWER 40 OF 57 MEDLINE L7
- Quantitation of metallothionein mRNA by RT-PCR and chemiluminescence. TΙ
- ANSWER 41 OF 57 MEDLINE L7

genes.

L7

TΤ

ANSWER 36 OF 57

- Is there a role for non-nucleoside reverse transcriptase inhibitors in the TItreatment of HIV infection?.
- ANSWER 42 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L7
- Comparative biological and biochemical evaluation of a diverse ΤI group of nonnucleoside reverse transcriptase inhibitors.
- DUPLICATE 29 ANSWER 43 OF 57 MEDLINE L7
- Biological and biochemical anti-HIV activity of the benzothiadiazine class TIof nonnucleoside reverse transcriptase inhibitors.
- DUPLICATE 30 ANSWER 44 OF 57 MEDLINE L7
- An insert of seven amino acids confers functional differences between TТ smooth muscle myosins from the intestines and vasculature.
- ANSWER 45 OF 57 CAPLUS COPYRIGHT 2002 ACS L7
- Use of a PCR-based method to characterize protein kinase C isoform ΤI expression in cardiac cells
- ANSWER 46 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. L7
- Variability in the reverse transcriptase TIgene, studied by direct DNA sequencing.
- ANSWER 47 OF 57 CAPLUS COPYRIGHT 2002 ACS L7
- HIV-1 reverse transcriptase: a diversity generator and quasispecies TIregulator
- MEDLINE DUPLICATE 31 ANSWER 48 OF 57 L7
- Comparison of HIV-1 and avian myeloblastosis virus reverse transcriptase TIfidelity on RNA and DNA templates.
- DUPLICATE 32 ANSWER 49 OF 57 MEDLINE L7
- Retroelements in bacteria. TΙ
- DUPLICATE 33 MEDLINE ANSWER 50 OF 57 L7
- Two independent retrons with highly diverse reverse ΤI transcriptases in Myxococcus xanthus.

- L7 ANSWER 51 OF 57 CAPLUS COPYRIGHT 2002 ACS
- TI Generation of diversity in retroviruses
- L7 ANSWER 52 OF 57 MEDLINE DUPLICATE 34
- TI Cell surface phenotype and human T lymphotropic virus type 1 antigen expression in 12 T cell lines derived from peripheral blood and cerebrospinal fluid of West Indian, Guyanese and African patients with tropical spastic paraparesis.
- L7 ANSWER 53 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI PHYLOGENETIC EVIDENCE FOR THE TRANSFER OF CASEOBACTER-POLYMORPHUS CROMBACH TO THE GENUS CORYNEBACTERIUM.
- L7 ANSWER 54 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI HUMAN T LYMPHOTROPIC VIRUSES AND DISEASES OF MAN.
- L7 ANSWER 55 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI MOLECULAR CLONING OF 7 MOUSE IMMUNO GLOBULIN K CHAIN MESSENGER RNA.
- L7 ANSWER 56 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
- TI NONCODING NUCLEOTIDE SEQUENCE IN THE 3-PRIME TERMINAL REGION OF A MOUSE IMMUNO GLOBULIN KAPPA CHAIN MESSENGER RNA DETERMINED BY ANALYSIS OF COMPLEMENTARY DNA.
- L7 ANSWER 57 OF 57 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI DEMONSTRATION THAT A MOUSE IMMUNO GLOBULIN LIGHT CHAIN MESSENGER RNA HYBRIDIZES EXCLUSIVELY WITH UNIQUE DNA.

=> s mmlv

L8 275 MMLV

=> s m(w) mlv

L9 140 M(W) MLV

=> s 18 or 19

L10 413 L8 OR L9

=> s 110 (9a) reverse (w) transcript? L11 133 L10 (9A) REVERSE (W) TRANSCRIPT?

=> dup rem 111

PROCESSING COMPLETED FOR L11

L12 75 DUP REM L11 (58 DUPLICATES REMOVED)

=> d 1-75 ti

- L12 ANSWER 1 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Inhibition of RNase using RNA heteropolymer in reverse transcription reaction
- L12 ANSWER 2 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI A novel gene organization: Intronic snoRNA gene clusters from Oryza sativa.
- L12 ANSWER 3 OF 75 MEDLINE DUPLICATE 1
- TI The role of template-primer in protection of reverse transcriptase from thermal inactivation.
- L12 ANSWER 4 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

- Base pairing properties of 8-oxo-7,8-dihydroadenosine in cDNA synthesis by TΙ reverse transcriptases. DUPLICATE 2 L12 ANSWER 5 OF 75 MEDLINE Low efficiency of the Moloney murine leukemia virus reverse transcriptase during reverse transcription of rare t(8;21) fusion gene transcripts. L12 ANSWER 6 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. Low efficiency of the Moloney murine leukemia virus reverse transcriptase during reverse transcription of rare t(8;21) fusion gene transcripts. DUPLICATE 3 L12 ANSWER 7 OF 75 MEDLINE Transcriptional profiling of a human papillomavirus 33-positive squamous TΙ epithelial cell line which acquired a selective growth advantage after viral integration.
- L12 ANSWER 8 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Evidence that BmTXK.beta.-BmKCT cDNA from Chinese scorpion Buthus martensii Karsch is an artifact generated in the reverse transcription process
- L12 ANSWER 9 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Localization of transcripts corresponding to the major allergen from olive pollen (Ole e I) by electron microscopic non-radioactive in situ RT-PCR.
- L12 ANSWER 10 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI GB virus C infection in blood donors from Cordoba, Argentina.
- L12 ANSWER 11 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Modified or mutated reverse transcriptases with high thermostability and uses thereof
- L12 ANSWER 12 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI High fidelity reverse transcriptases which have been modified or mutated and uses thereof
- L12 ANSWER 13 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI one step RT-PCR methods using enzyme mixes and kits comprising mutant thermostable polymerase and reverse transcriptase
- L12 ANSWER 14 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Improving reverse transcription at high temperatures using thermostable CpkB Chaperonin from hyperthermophilic archaeon Pyrococcus
- L12 ANSWER 15 OF 75 MEDLINE DUPLICATE 4
- TI Detection of the 5'-cap structure of messenger RNAs with the use of the cap-jumping approach.
- L12 ANSWER 16 OF 75 MEDLINE DUPLICATE 5
- TI Reverse transcriptase incorporation of 1,5-anhydrohexitol nucleotides.
- L12 ANSWER 17 OF 75 MEDLINE DUPLICATE 6
- TI A directed approach to improving the solubility of Moloney murine leukemia virus reverse transcriptase.
- L12 ANSWER 18 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Structure of a pseudo-16-mer DNA with stacked guanines and two G-A mispairs complexed with the N-terminal fragment of Moloney murine leukemia virus reverse transcriptase
- L12 ANSWER 19 OF 75 MEDLINE DUPLICATE 7
- TI Reverse transcriptase template switching: a SMART approach for full-length

cDNA library construction.

- L12 ANSWER 20 OF 75 MEDLINE DUPLICATE 8
- TI Construction of cDNA library of Eimeria tenella sporulated oocysts.
- L12 ANSWER 21 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Construction of cDNA library of Epinephelus cpoioies leukocytes
- L12 ANSWER 22 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Cytokine gene expression microarrays in the Rhesus model of Lyme neuroborreliosis.
- L12 ANSWER 23 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Comparative expression profiles of ETV6, CBFA2 and ETV6-CBFA2 in disease and remission states in childhood acute leukemia.
- L12 ANSWER 24 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Mutant form reverse transcriptase of Moloney murine leukemia virus with improved reactivity at high temperature
- L12 ANSWER 25 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI In situ hybrdization for RNA: Nonradioactive probe: ss cDNA probe.
- L12 ANSWER 26 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Analysis of plus-strand primer selection, removal, and reutilization by retroviral reverse transcriptases
- L12 ANSWER 27 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI One-step RT-PCR for detection of bluetongue virus RNA
- L12 ANSWER 28 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Construction of oocyte cDNA libraries of gynogenetic silver crucian carp and gonochoristic color crucian carp and cloning of their cyclin Al cDNAs
- L12 ANSWER 29 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Exonuclease III-generated series of homologous competitor DNA fragments for competitive PCR.
- L12 ANSWER 30 OF 75 CAPLUS COPYRIGHT 2002 ACS
- Reverse transcription of a naturally occurring nonretroviral RNA produces a precise deletion in the majority of its cDNA products
- L12 ANSWER 31 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Identification of age-associated genes in rat and mice brain by differential display PCR with selected primers.
- L12 ANSWER 32 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Analysis of gene expression following spinal cord injury using cDNA microarray technology.
- L12 ANSWER 33 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Insertional RNA editing in metazoan mitochondria: The cytochrome b gene in the nematode Teratocephalus lirellus.
- L12 ANSWER 34 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Update to: Automated recording of RNA differential display patterns from pig granulosa cells.
- L12 ANSWER 35 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE
- TI Misreading of RNA templates containing 8-oxo-7,8-dihydroguanosine or 8-oxo-2'-O-methylguanosine in cDNA synthesis by reverse transcriptases.

DUPLICATE 10 L12 ANSWER 36 OF 75 MEDLINE Molecular identification and immunolocalization of the water channel protein aquaporin 1 in CBCECs. L12 ANSWER 37 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.DUPLICATE Cloning and sequencing of the cDNA encoding for pokeweed anti-viral TΙ protein (PAP) and construction of its plant expression vector. L12 ANSWER 38 OF 75 CAPLUS COPYRIGHT 2002 ACS Molecular cloning of E-selectin from human umbilical vein endothelial cells DUPLICATE 12 L12 ANSWER 39 OF 75 MEDLINE Oligoribonucleotides containing 8-oxo-7,8-dihydroguanosine and

- Oligoribonucleotides containing 8-oxo-7,8-dihydroguanosine and 8-oxo-7,8-dihydro-2'-O-methylguanosine: synthesis and base pairing properties.
- L12 ANSWER 40 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Quantitative determination of cyclooxygenase-2 (COX-2) mRNA expression in peripheral blood leukocytes with RT-PCR and fluorescent dye capillary electrophoresis.
- L12 ANSWER 41 OF 75 MEDLINE DUPLICATE 13
- TI A sensitive and robust method for measles RNA detection.
- L12 ANSWER 42 OF 75 MEDLINE DUPLICATE 14
- TI Efficient in vitro inhibition of HIV-1 gag reverse transcription by peptide nucleic acid (PNA) at minimal ratios of PNA/RNA.
- L12 ANSWER 43 OF 75 MEDLINE DUPLICATE 15
- TI Synthesis of full-length potyvirus cDNA copies suitable for the analysis of genome polymorphism.
- L12 ANSWER 44 OF 75 MEDLINE DUPLICATE 16
- TI Detection of the induction of Salmonella enterotoxin gene expression by contact with epithelial cells with RT-PCR.
- L12 ANSWER 45 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI The type of reverse transcriptase affects the sensitivity of some reverse transcription PCR methods
- L12 ANSWER 46 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Representative cDNA synthesis from nanogram level of total RNA: A novel method using the template switching reaction catalyzed by M-MLV reverse transcriptase.
- L12 ANSWER 47 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Anti-HIV activities and mechanisms of antisense oligonucleotides
- L12 ANSWER 48 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Fidelity of MMLV reverse transcriptase and Thermus thermophilus DNA polymerase during reverse transcription and DNA amplification.
- L12 ANSWER 49 OF 75 MEDLINE
- TI Inhibition of gene expression by antisense DNA.
- L12 ANSWER 50 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI Use of 33P-labeled primer increases the sensitivity and specificity of mRNA differential display

- L12 ANSWER 51 OF 75 MEDLINE DUPLICATE 17
- TI Two different PCR assays to detect enteroviral RNA in CSF samples from patients with acute aseptic meningitis.
- L12 ANSWER 52 OF 75 MEDLINE DUPLICATE 18
- TI Detection of hepatitis C virus RNA by a reliable, optimized single-step reverse transcription polymerase chain reaction.
- L12 ANSWER 53 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- TI Fidelity of MMLV reverse transcriptase and Thermus thermophilus DNA polymerase during reverse transcription and DNA amplification.
- L12 ANSWER 54 OF 75 MEDLINE DUPLICATE 19
- TI Comparison of M-MLV reverse transcriptase and Tth polymerase activity in RT-PCR of samples with low virus burden.
- L12 ANSWER 55 OF 75 MEDLINE DUPLICATE 20
- TI [Analysis of effectiveness of cDNA synthesis, induced using complementary primers and primers containing a noncomplementary base matrix].

  Analiz effektivnost' sinteza kDNK, initsiirovannogo s komplementarnykh praimerov i praimerov, soderzhashchikh nekomplementarnye matritse osnovaniia.
- L12 ANSWER 56 OF 75 MEDLINE DUPLICATE 21
- TI [Expression of cytokines and interferon-related genes in the mouse embryo].

  Expression des genes des cytokines et des genes associes a l'interferon chez l'embryon de la souris.
- L12 ANSWER 57 OF 75 MEDLINE DUPLICATE 22
- TI Expression and role of c-myc protooncogene in murine preimplantation embryonic development.
- L12 ANSWER 58 OF 75 MEDLINE DUPLICATE 23
- TI Lactoferrin cDNA. Expression and in vitro mutagenesis.
- L12 ANSWER 59 OF 75 MEDLINE DUPLICATE 24
- [Derivatives of ddUTP, modified at the 5-position of uridine, as substrate terminators of reverse transcriptase. Hydrolysis of oligonucleotides, terminated by these analogs, by phosphodiesterase I]. Proizvodnye ddUTP, modifitsirovannye v 5-polozhenii uridina, kak substratnye terminatory obratnykh transkriptaz. Gidroliz oligonukleotidov, terminirovannykh etimi analogami, fosfodiesterazoi I.
- L12 ANSWER 60 OF 75 CAPLUS COPYRIGHT 2002 ACS
- TI A 'one tube reaction' for synthesis and amplification of total cDNA from small numbers of cells
- L12 ANSWER 61 OF 75 MEDLINE DUPLICATE 25
- TI [Expression of cytokine messenger RNA in murine placenta].
  Expression de l'ARN messager des cytokines dans le placenta de la souris.
- L12 ANSWER 62 OF 75 MEDLINE DUPLICATE 26
- [Reverse transcriptase of the human immunodeficiency virus: isolation and substrate specificity].

  Obratnaia transkriptaza virusa immunodefitsita cheloveka: vydelenie i substratnaia spetsifichenost'.
- L12 ANSWER 63 OF 75 MEDLINE

Ribosome initiation complex formation with the pseudoknotted alpha operon ΤI messenger RNA. DUPLICATE 28 L12 ANSWER 64 OF 75 MEDLINE [Induction of messenger RNA of cytokines by Herpes simplex virus infection TΙ Induction de l'ARN messager des cytokines par l'infection de l'Herpes simplex virus chez la souris. DUPLICATE 29 MEDLINE L12 ANSWER 65 OF 75 [Expression of cytokine messenger RNA in mice in physiological ΤI conditions]. Expression de l'ARN messager des cytokines chez la souris dans des conditions physiologiques. DUPLICATE 30 L12 ANSWER 66 OF 75 MEDLINE C-MYC mRNA is present in human sperm cells. TIMEDLINE DUPLICATE 31 L12 ANSWER 67 OF 75 Quantitation of changes in the expression of multiple genes by simultaneous polymerase chain reaction. DUPLICATE 32 L12 ANSWER 68 OF 75 MEDLINE Exogenous primer-independent cDNA synthesis with commercial reverse TТ transcriptase preparations on plant virus RNA templates. L12 ANSWER 69 OF 75 CAPLUS COPYRIGHT 2002 ACS Nucleotide sequence of a porcine prepro atrial natriuretic peptide (ANP) TI cDNA DUPLICATE 33 L12 ANSWER 70 OF 75 MEDLINE Low-ratio hybridization subtraction. DUPLICATE 34 L12 ANSWER 71 OF 75 MEDLINE Rapid amplification of complementary DNA from small amounts of TТ unfractionated RNA. DUPLICATE 35 L12 ANSWER 72 OF 75 MEDLINE Alpha-anomeric DNA: beta-RNA hybrids as new synthetic inhibitors of Escherichia coli RNase H, Drosophila embryo RNase H and  ${\bf M}$ -MLV reverse transcriptase. DUPLICATE 36 L12 ANSWER 73 OF 75 MEDLINE Isolation of cloned Moloney murine leukemia virus reverse transcriptase lacking ribonuclease H activity. DUPLICATE 37 L12 ANSWER 74 OF 75 MEDLINE Cloning and overexpression of Moloney murine leukemia virus reverse TТ transcriptase in Escherichia coli. L12 ANSWER 75 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC. Expression of cytokine and interferon-related genes in mouse embryo. => d 53 bib ab L12 ANSWER 53 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.

AN 1995:326610 BIOSIS

DN PREV199598340910

TI Fidelity of MMLV reverse transcriptase and Thermus thermophilus DNA polymerase during reverse transcription and DNA amplification.

- AU Myers, Thomas W.; Sigua, Christopher L.; Lawyer, Frances C.; Gelfand, David H.
- CS Program Core Res., Roche Molecular Systems, Alameda, CA 94501 USA
- Journal of Cellular Biochemistry Supplement, (1995) Vol. 0, No. 21A, pp. 302.

Meeting Info.: Keystone Symposium on Repair and Processing of DNA Damage Taos, New Mexico, USA March 23-29, 1995 ISSN: 0733-1959.

- DT Conference
- LA English

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- L12 ANSWER 48 OF 75 BIOSIS COPYRIGHT 2002 BIOLOGICAL ABSTRACTS INC.
- AN 1995:286429 BIOSIS
- DN PREV199598300729
- TI Fidelity of MMLV reverse transcriptase and Thermus thermophilus DNA polymerase during reverse transcription and DNA amplification.
- AU Sigua, Christopher L.; Lawyer, Frances C.; Gelfand, David H.; Myers, Thomas W.
- CS Program Core Res., Roche Molecular Systems, Alameda, CA 94501 USA
- FASEB Journal, (1995) Vol. 9, No. 6, pp. A1336.
  Meeting Info.: Annual Meeting of the American Society for Biochemistry and
  Molecular Biology San Francisco, California, USA May 21-25, 1995
  ISSN: 0892-6638.
- DT Conference
- LA English

### => d 24 45 bib ab

- L12 ANSWER 24 OF 75 CAPLUS COPYRIGHT 2002 ACS
- AN 2000:344167 CAPLUS
- DN 133:2042
- TI Mutant form reverse transcriptase of Moloney murine leukemia virus with improved reactivity at high temperature
- IN Arakawa, Taku; Nishiya, Yoshiaki; Kawakami, Fumikiyo; Kawamura, Yoshihisa
- PA Toyobo Co., Ltd., Japan
- SO Jpn. Kokai Tokkyo Koho, 10 pp. CODEN: JKXXAF
- DT Patent
- LA Japanese

### FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2000139457	A2	20000523	JP 1998-319241	19981110

PI JP 2000139457 A2 20000523 JP 1998-319241 19981110

AB A mutant form enzyme (V224M + D584N) of Moloney murine leukemia virus (

MMLV)-derived reverse transcriptase is

provided by point mutation so that the reactivity at a high temp. range (esp., extending ability at 42-60.degree.C) is improved comparing to the wild type and the conventional mutant, and the full-length cDNA is obtained. The mutant enzyme carries no substantial RNase H activity, and contains Tyr-Met-Asp-Asp sequence instead of Tyr-Val-Asp-Asp for the conserved region Tyr-X-Asp-Asp. A vector carrying the recombinant DNA encoding this mutant enzyme, and recombinant host cells (Escherichia coli) transformed using this vector, are claimed.

- L12 ANSWER 45 OF 75 CAPLUS COPYRIGHT 2002 ACS
- AN 1997:258097 CAPLUS
- DN 126:302153

TI The type of reverse transcriptase affects the sensitivity of some reverse transcription PCR methods

AU Barragan-Gonzalez, E.; Lopez-Guerrero, J. A.; Bolufer-Gilabert, P.; Sanz-Alonso, M.; De la Rubia-Comos, J.; Sempere-Talens, A.

CS Molecular Biology Lab., Dep. Clinical Biochem., Hospital Univ. La Fe, Valencia, 46009, Spain

SO Clinica Chimica Acta (1997), 260(1), 73-83 CODEN: CCATAR; ISSN: 0009-8981

PB Elsevier

DT Journal

LA English

AB A comparison of the efficacy of avian myelomatosis virus (AMV) vs. murine moloney leukemia virus (MMLV) reverse transcriptase in PCR mutation detection.

### => d 17 bib ab

L12 ANSWER 17 OF 75 MEDLINE

DUPLICATE 6

AN 2001520141 MEDLINE

DN 21451146 PubMed ID: 11567084

TI A directed approach to improving the solubility of Moloney murine leukemia virus reverse transcriptase.

AU Das D; Georgiadis M M

CS Waksman Institute and Department of Chemistry and Chemical Biology, Rutgers University, Piscataway, New Jersey 08854, USA.

NC GM 55026 (NIGMS)

SO PROTEIN SCIENCE, (2001 Oct) 10 (10) 1936-41. Journal code: 9211750. ISSN: 0961-8368.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200112

ED Entered STN: 20010924
Last Updated on STN: 20020122
Entered Medline: 20011205

One of the difficulties that can impede structural work on a molecule of AΒ interest is limited solubility. Although functionally similar to the human immunodeficiency virus type-1 reverse transcriptase (HIV-1 RT), the Moloney murine leukemia virus reverse transcriptase (MMLV RT) differs both in architecture and solubility properties. Reverse transcriptase is an essential retroviral enzyme that replicates the single-stranded RNA genome of the retrovirus producing a double-stranded DNA copy, which is subsequently integrated into the host's genome. We have introduced a single amino acid substitution in the connection domain of an N-terminally truncated MMLV RT (L435K) that significantly improves the solubility of the enzyme eliminating the need for nonionic detergents in buffering storage solutions. The substituted enzyme retains near wild-type polymerase activity. An important consequence of the improved solubility of the L435K MMLV RT has been the ability to obtain diffraction quality crystals.

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L12 ANSWER 73 OF 75 MEDLINE DUPLICATE 36

88124200 MEDLINE AN

PubMed ID: 2448747 88124200 DN

- Isolation of cloned Moloney murine leukemia virus reverse transcriptase TΤ lacking ribonuclease H activity.
- Kotewicz M L; Sampson C M; D'Alessio J M; Gerard G F ΑU
- Molecular Biology Research and Development, Bethesda Research CS Laboratories, Life Technologies, Inc., Gaithersburg, MD 20877.
- NUCLEIC ACIDS RESEARCH, (1988 Jan 11) 16 (1) 265-77. Journal code: 0411011. ISSN: 0305-1048. SO

ENGLAND: United Kingdom CY

- Journal; Article; (JOURNAL ARTICLE) DT
- English LΑ
- FS Priority Journals
- EM 198803
- Entered STN: 19900308 ED Last Updated on STN: 19970203 Entered Medline: 19880307
- Retroviral reverse transcriptase possesses DNA polymerase and ribonuclease AΒ H (RNase H) activity within a single polypeptide. Chemical or proteolytic treatment of reverse transcriptase has been used in the past to produce enzyme that is missing DNA polymerase activity and retains RNase H activity. It has not been possible to obtain reverse transcriptase that lacks RNase H but retains DNA polymerase activity. We have constructed a novel deletion derivative of the cloned Moloney murine leukemia virus ( M-MLV) reverse transcriptase gene,
- expressed the gene in E. coli, and purified the protein to near homogeneity. The purified enzyme has a fully active DNA polymerase, but has no detectable RNase H activity. These results are consistent with, but do not prove, the conclusion that the DNA polymerase and RNase H activities of M-MLV reverse

transcriptase reside within separate structural domains.

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L12 ANSWER 24 OF 75 CAPLUS COPYRIGHT 2002 ACS
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AN 2000:344167 CAPLUS

DN 133:2042

TI Mutant form reverse transcriptase of Moloney murine leukemia virus with improved reactivity at high temperature

IN Arakawa, Taku; Nishiya, Yoshiaki; Kawakami, Fumikiyo; Kawamura, Yoshihisa

PA Toyobo Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 10 pp. CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PI JP 2000139457 A2 20000523 JP 1998-319241 19981110 AB A mutant form enzyme (V224M + D584N) of Moloney murine leukemia virus (

MMLV)-derived reverse transcriptase is provided by point mutation so that the reactivity at a high temp. range (esp., extending ability at 42-60.degree.C) is improved comparing to the wild type and the conventional mutant, and the full-length cDNA is obtained. The mutant enzyme carries no substantial RNase H activity, and contains Tyr-Met-Asp-Asp sequence instead of Tyr-Val-Asp-Asp for the conserved region Tyr-X-Asp-Asp. A vector carrying the recombinant DNA encoding this mutant enzyme, and recombinant host cells (Escherichia coli) transformed using this vector, are claimed.